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(FILE 'HOME' ENTERED AT 14:11:00 ON 07 MAR 2003)

     FILE 'REGISTRY' ENTERED AT 14:11:10 ON 07 MAR 2003
                E "IMIDAZOLE-4-ACETIC ACID-RIBOSE"/CN 25
L1              STRUCTURE UPLOADED
L2            0 S L1
L3            0 S L1 SSS SAM
L4            5 S L1 SSS FULL

     FILE 'CAPLUS, MEDLINE, USPATFULL' ENTERED AT 14:22:59 ON 07 MAR 2003
L5           28 S L4
L6            0 S L5 AND IMIDAZOLINE
L7            0 S L5 AND IMIDAZOLINE
L8            0 S L5 AND CELL
L9            2 S L5 AND RECEPTOR
L10          26 S L5 AND IMIDAZOL?
L11           0 S L10 AND CONGENER
L12         111 S L0 AND (HYPERTENSION OR GLAUCOMA OR PSYCHIATRIC OR DIABETES O
L13           0 S LL0 AND (HYPERTENSION OR GLAUCOMA OR PSYCHIATRIC OR DIABETES
L14           0 S L10 AND CONTACT?
L15           0 S L10 AND ARACHIDONIC

=>
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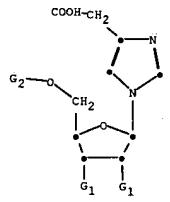
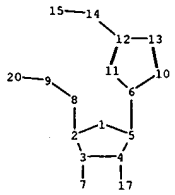




chain nodes :
7 8 9 14 15 17 20
ring nodes :
1 2 3 4 5 6 10 11 12 13
chain bonds :
2-8 3-7 4-17 5-6 8-9 9-20 12-14 14-15
ring bonds :
1-2 1-5 2-3 3-4 4-5 6-10 6-11 10-13 11-12 12-13
exact/norm bonds :
1-2 1-5 2-3 3-4 3-7 4-5 4-17 5-6 6-10 6-11 9-20 10-13 11-12 12-13
exact bonds :
2-8 8-9 12-14 14-15

G1:OH,X,Ak

G2:H,PO3H2

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:Atom
11:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 17:CLASS 20:Atom

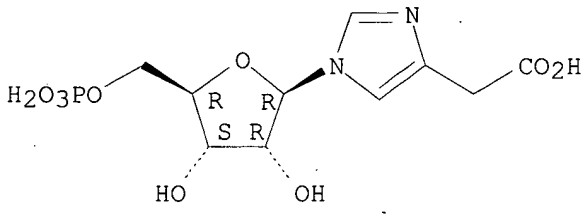
L10 ANSWER 1 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1995:715348 CAPLUS
DOCUMENT NUMBER: 123:103310
TITLE: **Imidazoleacetic** acid, a .gamma.-aminobutyric acid receptor agonist, can be formed in rat brain by oxidation of histamine
AUTHOR(S): Thomas, Boban; Prell, George D.
CORPORATE SOURCE: Dep. Pharmacol., Mt. Sinai Sch. Med. City Univ. New York, New York, NY, USA
SOURCE: Journal of Neurochemistry (1995), 65(2), 818-26
CODEN: JONRA9; ISSN: 0022-3042
PUBLISHER: Lippincott-Raven
DOCUMENT TYPE: Journal
LANGUAGE: English

AB It is generally accepted that in mammalian brain histamine is metabolized solely by histamine methyltransferase (HMT), to form tele-methylhistamine, then oxidized to tele-methylimidazoleacetic acid. However, histamine's oxidative metabolite in the periphery, **imidazoleacetic** acid (IAA), is also present in brain and CSF, and its levels in brain increase after inhibition of HMT. To reinvestigate if brain has the capacity to oxidize histamine and form IAA, conscious rats were injected with [3H]histamine (10 ng), either into the lateral ventricles or cisterna magna, and decapitated 30 min later. In brains of saline-treated rats, most radioactivity recovered was due to tele-methylhistamine and tele-methylimidazoleacetic acid. However, significant amts. of tritiated IAA and its metabolites, IAA-ribotide and IAA-riboside, were consistently recovered. In rats pretreated with metoprine, an inhibitor of HMT, labeled IAA and its metabolites usually comprised the majority of histamine's tritiated metabolites. [3H]Histamine given intracisternally produced only trace amts. of oxidative metabolites. Formation of IAA, a potent GABA-A agonist with numerous neurochem. and behavioral effects, from minute quantities of histamine in brain indicates a need for reevaluation of histamine's metabolic pathway or pathways in brain and suggests a novel mechanism for interactions between histamine and the GABAergic system.

IT **2888-19-9, Imidazole**-4-acetic acid-ribotide
29605-99-0, Imidazoleacetic acid-riboside
RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL (Biological study); FORM (Formation, nonpreparative)
(**imidazoleacetic** acid formation in rat brain by histamine oxidn.)

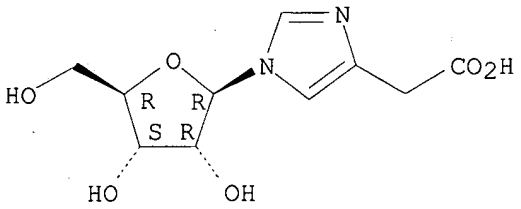
RN 2888-19-9 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

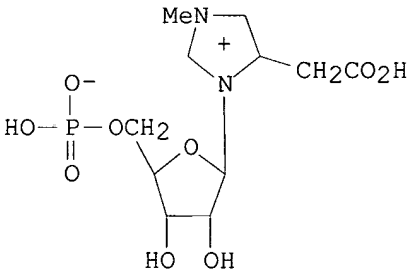


RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 2 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1992:470204 CAPLUS
DOCUMENT NUMBER: 117:70204
TITLE: Nucleosides. 163. Synthesis of ribosides and ribotides of **imidazole**-4(5)-acetic acid and 1-methylimidazole-4(5)-acetic acid
AUTHOR(S): Matulic-Adamic, Jasenka; Watanabe, Kyoichi A.
CORPORATE SOURCE: Lab. Org. Chem., Sloan-Kettering Inst. Cancer Res., New York, NY, 10021, USA
SOURCE: Korean Journal of Medicinal Chemistry (1991), 1(1), 54-64
CODEN: KJMCE7; ISSN: 1225-0058
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 117:70204
GI

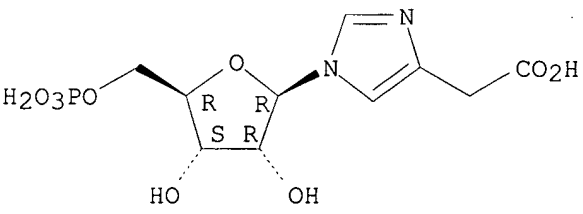


I

AB Nucleotide **imidazoleacetic** acid, e.g. I, were prepd. from **imidazole**-4(5)-acetonitrile (II). Regioselective tritylation of II followed by N-methylation with Me2S and hydrolysis gave 1-methylimidazole-5-acetic acid.

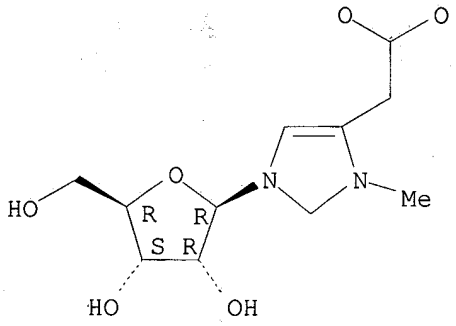
IT **2888-19-9P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and N-methylation of)
RN 2888-19-9 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

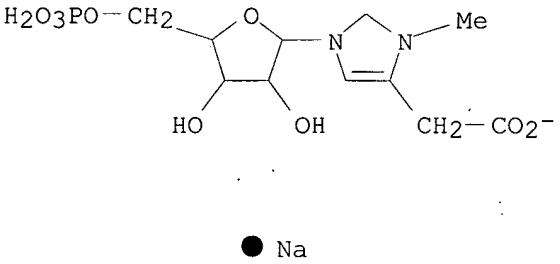


IT **142527-55-7P 142606-76-6P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)
RN 142527-55-7 CAPLUS
CN 1H-Imidazolium, 4-(carboxymethyl)-3-methyl-1-.beta.-D-ribofuranosyl-, inner salt (9CI) (CA INDEX NAME)

Absolute stereochemistry.

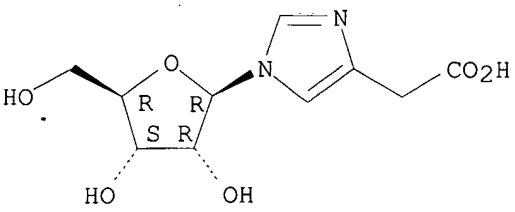


*** FRAGMENT DIAGRAM IS INCOMPLETE ***
RN 142606-76-6 CAPLUS
CN 1H-Imidazolium, 4-(carboxymethyl)-3-methyl-1-(5-O-phosphono-.beta.-D-ribofuranosyl)-, inner salt, monosodium salt (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***
IT **29605-99-0P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn., N-methylation, and phosphorylation of)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 3 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1984:18260 CAPLUS
DOCUMENT NUMBER: 100:18260
TITLE: Catabolism of histamine in the isolated glomeruli and

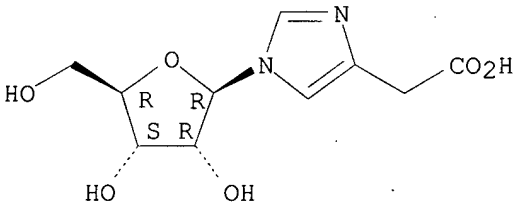
tubules of the rat kidney
AUTHOR(S): Abboud, Hanna E.
CORPORATE SOURCE: Dep. Med., Case West. Res. Univ., Cleveland, OH, USA
SOURCE: Kidney International (1983), 24(4), 534-41
CODEN: KDYIA5; ISSN: 0085-2538
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Rat kidney glomeruli and cortical tubules were incubated with radiolabeled histamine [51-45-6], and the products were sepd. by TLC. Glomeruli predominantly catabolized histamine to acid metabolites of the diamine oxidase (histaminase) pathway, **imidazole** acetic acid [645-65-8] and ribosylimidazole acetic acid [**29605-99-0**], and to a lesser extent to the inactive methylation product, N.tau.-methylhistamine [501-75-7]. Tubules, on the other hand, catabolized histamine to N.tau.-methylhistamine and to a lesser degree to acid metabolites. The Me donor S-adenosyl-methionine (SAM) (10-4M) markedly enhanced the prodn. of N.tau.-methylhistamine in both glomeruli and tubules but had no effect on the prodn. of acid metabolites. In the presence of equimolar concns. of SAM, tubules continued to methylate histamine to a greater extent than glomeruli. In both glomeruli and tubules, the diamine oxidase inhibitor, amino-guanidine, abolished the prodn. of acid metabolites whereas amodiaquine and pyrilamine, inhibitors of the methylation pathway, markedly reduced the prodn. of N.tau.-methylhistamine. In the presence of SAM, tubules catabolized nonlabeled histamine to a greater extent than glomeruli. Thus, tubules have a greater capacity than glomeruli to degrade histamine and histamine is differentially catabolized in these segments. A major pathway of histamine catabolism in glomeruli results in the formation of biol. active products.

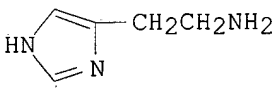
IT **29605-99-0**
RL: BIOL (Biological study)
(as histamine metabolite, in kidney glomeruli)

RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 4 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1983:138012 CAPLUS
DOCUMENT NUMBER: 98:138012
TITLE: Biliar elimination of histamine and its metabolites in guinea pigs
AUTHOR(S): Puerta, M. L.; Ballestero, M. E. M.
CORPORATE SOURCE: Fac. Cienc. Biol., Univ. Complutense Madrid, Madrid, Spain
SOURCE: Comparative Biochemistry and Physiology, C: Comparative Pharmacology (1983), 74C(1), 111-13
CODEN: CBPCBB; ISSN: 0306-4492
DOCUMENT TYPE: Journal
LANGUAGE: English
GI



I

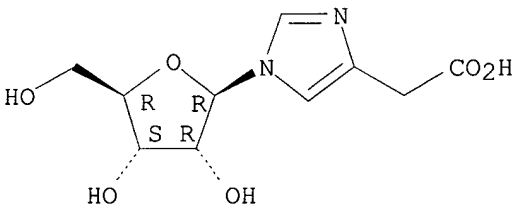
AB Administration of 14C-labeled histamine (I) [51-45-6] i.v. to guinea pigs resulted in 3.5% of the radioactivity being eliminated in the bile of both males and females. Free I, methylhistamine [501-75-7], methimidazoleacetic acid [2625-49-2], **imidazoleacetic** acid [645-65-8] and its riboside [**29605-99-0**], and acetylhistamine [673-49-4] were identified in the bile. Male bile contained more free I and methylhistamine than did female bile. Evidently, biliary elimination of I and metabolites is similar to that of urine but quant. less important.

IT **29605-99-0**
RL: BIOL (Biological study)
(as histamine metabolite, in bile)

RN 29605-99-0 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 5 OF 26 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1978:48665 CAPLUS

DOCUMENT NUMBER: 88:48665

TITLE: Histamine metabolism in cluster headache and migraine. Catabolism of 14C-histamine

AUTHOR(S): Sjaastad, Ottar; Sjaastad, O. V.

CORPORATE SOURCE: Dep. Neurol., Rikshosp., Oslo, Norway

SOURCE: Journal of Neurology (1977), 216(2), 105-17
CODEN: JNRYA9; ISSN: 0340-5354

DOCUMENT TYPE: Journal

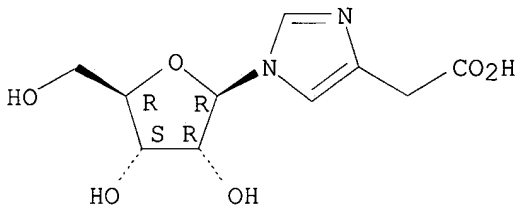
LANGUAGE: English

AB Various parameters of histamine metab. were studied in patients with migraine, cluster headache, and chronic paroxysmal hemicrania. These included urinary excretion of radioactivity and of histamine-14C and its metabolites, exhaled 14CO2 and fecal radioactivity after oral as well as s.c. administration of histamine-14C. No marked deviation from the normal was found except in 1 patient with the cluster headache variant, chronic paroxysmal hemicrania, in whom an aberration in histamine degrdn. seemed to be present. Only min. quantities of the histamine-14C metabolite **imidazoleacetic**-14C acid riboside seemed to be formed during severe paroxysms. During a symptom-free period no deviation from normal was obsd. The most likely explanation for this finding seems to be a defect in the conversion of **imidazoleacetic** acid to its riboside. This defect may possibly explain the increased urinary excretion of histamine in this particular patient. The relation of this metabolic aberration to the prodn. of headache still remains dubious for various reasons.

IT **29605-99-0**
RL: FORM (Formation, nonpreparative)

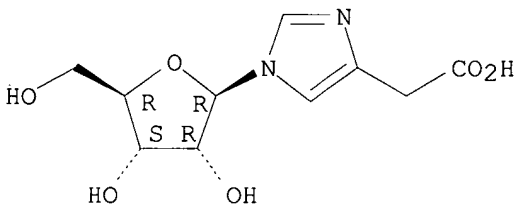
(formation of, from histamine during headache)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 6 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1977:101217 CAPLUS
DOCUMENT NUMBER: 86:101217
TITLE: Histamine and its metabolites in cat portal venous blood and intestine after duodenal instillation of histamine
AUTHOR(S): Marley, E.; Thomas, D. V.
CORPORATE SOURCE: Dep. Pharmacol., Inst. Psychiatry, London, UK
SOURCE: Journal of Physiology (Cambridge, United Kingdom) (1976), 263(2), 273P-274P
CODEN: JPHYA7; ISSN: 0022-3751
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Histamine [51-45-6] (0.175-0.220 mg/kg) instilled into the intestine of cats was metabolized mainly in the intestinal wall by either diamine oxidase [9001-53-0]-deamination or by **imidazole** N-methylation followed by monoamine oxidase [9001-66-5]-deamination. I.v. infused histamine (0.15 .mu.g/kg/min) was metabolized mainly by methylation at sites other than the intestine. Deamination by diamine oxidase was delayed in the absence of the gut and liver.
IT **29605-99-0**
RL: BIOL (Biological study)
(as histamine metabolite in intestine)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 7 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1976:553943 CAPLUS
DOCUMENT NUMBER: 85:153943
TITLE: Interference with histamine and **imidazole** acetic acid metabolism by salicylates: a possible contribution to salicylate analgesic activity?
AUTHOR(S): Beaven, M. A.; Horakova, Zdenka; Keiser, H. R.
CORPORATE SOURCE: Natl. Heart Lung Inst., NIH, Bethesda, MD, USA

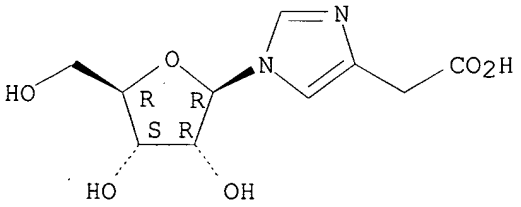
SOURCE: Experientia (1976), 32(9), 1180-2
CODEN: EXPEAM; ISSN: 0014-4754
DOCUMENT TYPE: Journal
LANGUAGE: English

AB In man, rats and mice, the urinary excretion of the histamine [51-45-6] and the L-histidine [71-00-1] metabolite, **imidazole** acetic acid [645-65-8], was increased and that of the conjugated metabolite, ribosylimidazole acetic acid [**29605-99-0**], decreased by small doses of salicylates. In contrast to salicylates, other non-salicylate anti-inflammatory drugs, indomethacin [53-86-1], phenylbutazone [50-33-9], phenacetin [62-44-2] and acetaminophen [103-90-2] did not influence the excretion of the urinary metabolites of histamine and L-histidine. Since **imidazole** acetic acid is reported to have analgesic and narcotic activity, there is the inference that the analgesic properties of salicylate might be due in part to interference in **imidazole** acetic acid metab.

IT **29605-99-0**
RL: BIOL (Biological study)
(as histidine metabolite, salicylate effect on, analgesic activity in relation to)

RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



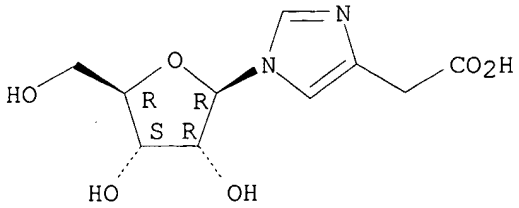
L10 ANSWER 8 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1975:166843 CAPLUS
DOCUMENT NUMBER: 82:166843
TITLE: Ion exchange chromatography for quantitative analysis of radioactive histamine metabolites in human urine
AUTHOR(S): Bergmark, J.; Granerus, G.
CORPORATE SOURCE: Dep. Clin. Chem., Univ. Goteborg, Goteborg, Swed.
SOURCE: Scandinavian Journal of Clinical and Laboratory Investigation (1974), 34(4), 365-73
CODEN: SJCLAY; ISSN: 0036-5513
DOCUMENT TYPE: Journal
LANGUAGE: English

AB The method was compared with the isotope diln. method developed by R. W. Schayer (1959). The advantages of the new method are rapidity and a greater possibility to account for all the radioactive histamine metabolites excreted in the urine. Data also suggested that the histamine metabolite, **imidazoleacetic** acid riboside, is partly lost in the isotope diln. method because of adsorption to urinary constituents during hydrolysis. The catabolism of i.v. and orally given histamine-14C was studied in one healthy subject. The general metabolic pattern of histamine was confirmed. In addn., it was found that histaminol might be a minor metabolite of injected histamine.

IT **29605-99-0**
RL: ANT (Analyte); ANST (Analytical study)
(detn. of, in urine as histamine metabolite)

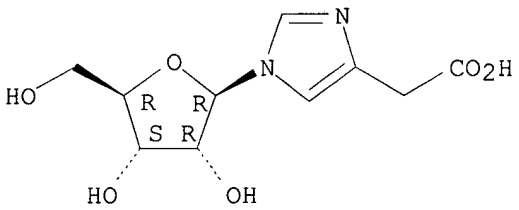
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 9 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1974:115938 CAPLUS
DOCUMENT NUMBER: 80:115938
TITLE: Catabolism of orally administered carbon-14-labeled histamine in man
AUTHOR(S): Sjaastad, Ottar; Sjaastad, O. V.
CORPORATE SOURCE: Inst. Surg. Res., Univ. Hosp., Oslo, Norway
SOURCE: Acta Pharmacologica et Toxicologica (1974), 34(1), 33-45
CODEN: APTOA6; ISSN: 0001-6683
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Within 48 hr following oral administration of 14C-labeled histamine-2HCl (I-2HCl) [56-92-8] (.sim.200 mg) to humans, 68-80% of the radioactivity was recovered in the urine, 1.8-18% was exhaled as 14CO2, and 13-19% was excreted in the feces. The main urinary I metabolites were **imidazoleacetic** acid [645-65-8] and methylimidazoleacetic acid [2625-49-2].
IT **29605-99-0**
RL: FORM (Formation, nonpreparative)
(formation of, as histamine metabolite)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

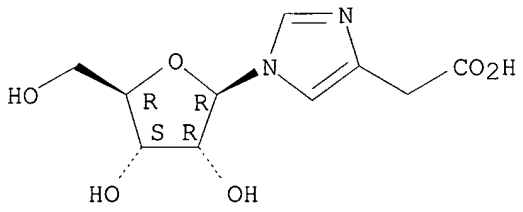


L10 ANSWER 10 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1973:146963 CAPLUS
DOCUMENT NUMBER: 78:146963
TITLE: Differentiation of 1,4- and 1,5-disubstituted **imidazoles**
AUTHOR(S): Matthews, H. Randall; Rapoport, Henry
CORPORATE SOURCE: Lawrence Berkeley Lab., Univ. California, Berkeley, CA, USA
SOURCE: Journal of the American Chemical Society (1973), 95(7), 2297-303
CODEN: JACSAT; ISSN: 0002-7863
DOCUMENT TYPE: Journal
LANGUAGE: English
AB A method is presented for distinguishing 1,4- and 1,5-disubstituted

imidazoles by their proton cross-ring coupling consts. Other spectral methods also have been evaluated, as well as several methods which have been reported for differentiating such isomers. Comparison of these methods leads to the conclusion that the measurement of cross-ring coupling consts. is the most generally satisfactory and reliable procedure. On this basis, structures are assigned to the carboxymethylhistidines, and the histamine metabolite is established as 1-.beta.-d-ribofuranosyl-4- **imidazoleacetic** acid.

IT **29605-99-0**
RL: PRP (Properties)
(NMR of)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

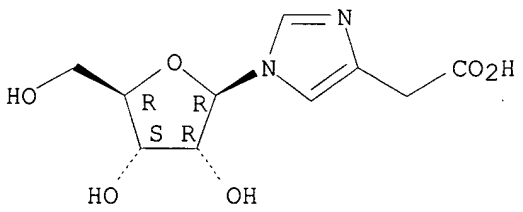
Absolute stereochemistry.



L10 ANSWER 11 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1971:137653 CAPLUS
DOCUMENT NUMBER: 74:137653
TITLE: Metabolism of [14C]-histamine in domestric animals. II. Cow and sheep
AUTHOR(S): Eliassen, K. A.
CORPORATE SOURCE: Dep. Physiol., Vet. Coll. Norway, Oslo, Norway
SOURCE: Acta Physiologica Scandinavica (1971), 81(3), 289-99
CODEN: APSCAX; ISSN: 0001-6772
DOCUMENT TYPE: Journal
LANGUAGE: English
GI For diagram(s), see printed CA Issue.
AB In cow and sheep the oxidative deamination of histamine (I) into **imidazoleacetic** acid and its riboside was the major metabolic pathway. About 2% of the urinary radioactivity following 14C-labeled I injection was due to histaminol.

IT **29605-99-0**
RL: FORM (Formation, nonpreparative)
(formation of, from histamine, by ruminants)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

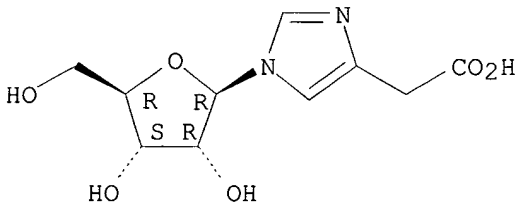
Absolute stereochemistry.



L10 ANSWER 12 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1971:74587 CAPLUS

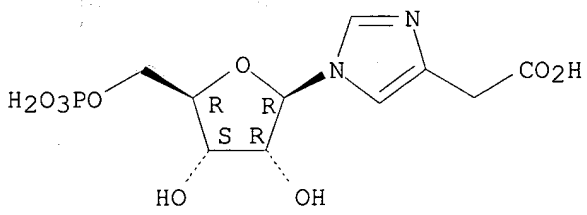
DOCUMENT NUMBER: 74:74587
TITLE: Uptake of [14C]-histamine by tissues of the guinea pig
AUTHOR(S): Lewis, A. J.; Nicholls, Paul J.
CORPORATE SOURCE: Welsh Sch. Pharm., UWIST, Cardiff, UK
SOURCE: Journal of Pharmacy and Pharmacology (1971), 23(1), 66
CODEN: JPPMAB; ISSN: 0022-3573
DOCUMENT TYPE: Journal
LANGUAGE: English
GI For diagram(s), see printed CA Issue.
AB The low uptake of ring-2-14C-labeled histamine (I) (80 mg/kg, i.v.) by various tissues of guinea pigs showed that the animal, unlike cats and rabbits, does not posses an effective uptake system. The acidic metabolites of I, when detd. 8 hr after the administration, were identified as **imidazole**-4-acetic acid, 1-ribosylimidazole-4-acetic acid, and 1-methylimidazole-4-acetic acid.
IT **29605-99-0**
RL: FORM (Formation, nonpreparative)
(formation of, from histamine by animal tissue)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 13 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1965:441136 CAPLUS
DOCUMENT NUMBER: 63:41136
ORIGINAL REFERENCE NO.: 63:7422g-h
TITLE: Evidence for the presence of **imidazoleacetic** acid riboside and ribotide in rat tissues
AUTHOR(S): Robinson, Joseph D.; Green, Jack P.
CORPORATE SOURCE: School of Med., Yale Univ.
SOURCE: Federation Proc. (1965), 24(3;1), 777
DOCUMENT TYPE: Journal
LANGUAGE: English
AB A combination of ion-exchange and paper chromatography of the acid-sol. radioactive material from kidneys of rats given histamine-14 C showed the presence of **imidazolcacetic** acid riboside (I) and ribotide (II) and a third unidentified substance whose Rf value differed from all known metabolites of histamine. The most likely route for the synthesis of I and II would be oxidn. of histamine to **imidazoleacetic** acid followed by condensation of the acid with phosphoribosyl pyrophosphate, a reaction demonstrated in vitro; the I would then arise by dephosphorylation. Labeled histamine adenine dinucleotide and histamine adenine dinucleotide phosphate could not be detected in kidney, liver, or brain.
IT **2888-19-9**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-phosphate
(in kidneys after histamine administration)
RN 2888-19-9 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



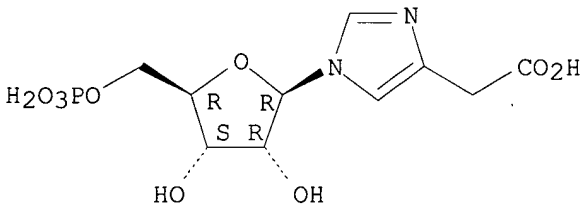
L10 ANSWER 14 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1965:441135 CAPLUS
DOCUMENT NUMBER: 63:41135
ORIGINAL REFERENCE NO.: 63:7422e-g
TITLE: Methylhistamine in urine and brain
AUTHOR(S): Fram, D. H.; Green, J. P.
CORPORATE SOURCE: School of Medicine, Yale Univ.
SOURCE: Federation Proc. (1965), 24(3;1), 778
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Methylhistamine (1-methyl-4-(.beta.-amino-ethyl)-**imidazole**) (I), found as a normal const. of urine and brain, was measured by prepg. its deriv. with 1-fluoro-2,4-dinitro-benzene (II) from acid exts. of urine or brain. The method was validated qual. by submitting the acid ext. to paper, thin-layer, ion-exchange, or gas chromatography and by submitting derivs. prepd. from the exts. with II, 2-fluoro-3-bromo-4,6-dinitro-benzene, 1-chloro-2,4,6-trinitrobenzene, or 2,4-dichloro-5-nitro-pyrimidine to paper or thin-layer chromatography. The 24-hr. urinary excretion of I in normal humans was 137-480 .gamma.; histamine excretion was 16-53 .gamma.. The ratio of I to histamine excreted ranged from 6 to 15. There was no sex difference in the excretion of either amine or in the ratios. The concn. of I in the brain of guinea pig was 45-75 -.gamma./g.

IT **2888-19-9**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-phosphate **29605-99-0**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-
(in kidneys after histamine administration)

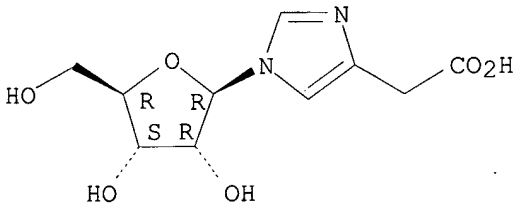
RN 2888-19-9 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 15 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1964:486555 CAPLUS
DOCUMENT NUMBER: 61:86555
ORIGINAL REFERENCE NO.: 61:15113g-h
TITLE: Presence of **imidazoleacetic** acid riboside and ribotide in rat tissues
AUTHOR(S): Robinson, J. D.; Green, J. P.
CORPORATE SOURCE: Yale Univ., School of Med., New Haven, CT
SOURCE: Nature (1964), 203(4950), 1178-9
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable

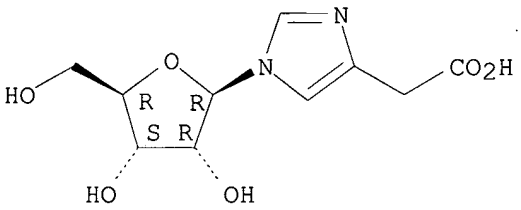
AB In rats given multiple injections of labeled histamine (I), chromatography of trichloroacetic acid (TCA) exts. of kidney revealed 3 major radioactive fractions. These were **imidazoleacetic** acid riboside (II), **imidazoleacetic** acid ribotide (III) and an unidentified fraction not coinciding with any of the urinary I metabolites. In brain, after injection of labeled histidine (IV), chromatography of TCA exts. revealed small fractions of total tissue radioactivity in II, III, I and **imidazoleacetic** acid, higher levels in unidentified metabolites, and at least 80% as IV. Radioactivity from injected I or IV was not incorporated into histamine adenine dinucleotide (HAD) or HAD phosphate in rat or guinea pig organs.

IT **29605-99-0**, **Imidazole**-4-acetic acid, 1-ribosyl-
(in brain and kidneys after administration of histamine and histidine)

RN 29605-99-0 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 16 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1964:471247 CAPLUS
DOCUMENT NUMBER: 61:71247
ORIGINAL REFERENCE NO.: 61:12423e-h
TITLE: The fate of histamine-14C in animal tissues
AUTHOR(S): Snyder, Solomon H.; Axelrod, Julius; Bauer, Hugo
CORPORATE SOURCE: Natl. Insts. of Health, Bethesda, MD
SOURCE: J. Pharmacol. Exptl. Therap. (1964), 144(3), 373-9
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable

AB Adult mice were administered histamine-2-14C (I) intravenously. There was a rapid initial disappearance of I. Methylhistamine-14C (II) concn. was max. at 30 min. After 90 min. I and II disappeared at a slower rate but could be detected in the whole animal at 48 hrs. After 48 hrs. the total

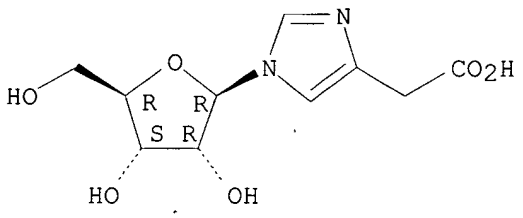
radioactivity was twice the sum of I and II. After 30 min. the concn. of II was the same or higher than that of I in most tissues. The ratio of II to I was greatest in the spleen. I was not found in the brain. In all tissues total radioactivity exceeded the sum of I and II. The ratio of total radioactivity to the sum of I and II was greatest in the liver and kidney, and least in the spleen, blood, skeletal muscle, and stomach. Liver and kidney had the greatest concn. of total activity. Adult rats of both sexes received I subcutaneously and were killed 1 or 24 hrs. later. Negligible amts. of II were found. In all tissues total activity exceeded the concn. of I at 1 and 24 hrs. At both times total activity was greatest in the kidney and the ratio to I was larger than in other tissues. After 1 hr. kidney and heart had the greatest concn. of I while brain, serum, and testes had the least. After 24 hrs. the spleen had the largest amt. of I and its ratio of total activity to I was lowest. There were detectable levels of I and total activity in serum after 24 hrs. **Imidazoleacetic** acid (III) and **imidazoleacetic** acid riboside (IV) were found in equal concns. At 90 min. and at 4 hrs. their concns. exceeded those of I and II. After 48 hrs. the sum of I and II equalled the sum of III and IV. After 1 hr. there was a larger amt. of III in heart, lung, intestine, and spleen; the kidney and liver had a greater concn. of IV. After 24 hrs. the above tissues contained 10 times as much IV as III. Methylimidazoleacetic acid-14 C was not found. The amts. of I found in the brain of the rat indicated that I would cross the blood-brain barrier.

IT **29605-99-0, Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-
(formation in histamine metabolism in tissues)

RN 29605-99-0 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 17 OF 26 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1964:456802 CAPLUS

DOCUMENT NUMBER: 61:56802

ORIGINAL REFERENCE NO.: 61:9880f-h,9881a

TITLE: Increased turnover of phosphoribosylpyrophosphate, a purine nucleotide precursor, in certain gouty subjects

AUTHOR(S): Wyngaarden, J. B.; Jones, O. W.; Ashton, D. M.

SOURCE: Atti Congr. Lega Intern. Reumatismo, 10.degree., Rome (1961), 1, 249-53

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Since phosphoribosylpyrophosphate (I) is an obligatory precursor of purine nucleotides, its turnover has been investigated in gouty subjects. The hyperuricemia of gout may be due to overproduction or underexcretion of uric acid, or both. Orally administered **imidazoleacetic** acid (II) is partially excreted in urine as the **imidazoleacetic** acid ribonucleotide (III), and if glucose-14C is given simultaneously the ribose moiety is labeled. It is assumed that the same "pool" of I is involved both in the production of III and of phosphoribosylamine (the 1st specific precursor of purine nucleotides). Subjects were all males. Five controls had no gout or renal disease personally, or in the family

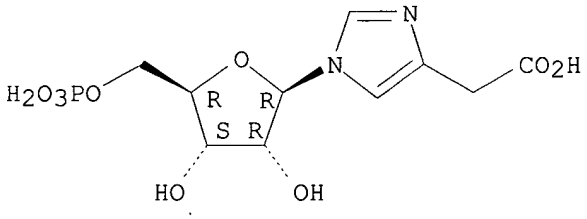
history. Five gouty patients varied from asymptomatic hyperuricemia to advanced chronic tophaceous gout. All were given 25 .mu.c. glucose-U-14C and 20 micromoles/kg. II. Urine was collected in 5 ml. of concd. HCl, either in 2-hr. aliquots, or in a single 10-hr. sample, and stored at 4.degree.. CO2 was then removed by aeration, the pH adjusted to 8, the III collected on a Dowex-1 (acetate) column, and purified on a Dowex-50 (H+) column. The product in M citrate, pH 6.0, was hydrolyzed with a bacterial riboside hydrolase, and the protein-free filtrate passed through a mixed-bed resin (MB-3, Fisher), and the eluate analyzed for 14C and ribose (orcinol). Uric acid was detd. by differential spectrophotometry using uricase. In the controls, 0.010-0.047% 14C was incorporated into urinary III in 10 hours. For 2 gouty subjects with low and normal uric acid excretions, the corresponding figures were 0.009 and 0.058%, and for 3 gouty hyperuricemic subjects the range was 0.1640.309%. In these latter 3 subjects, the sp. activity (counts/ min./mg.) of the ribose moiety of III was approx. 8 times that of the controls. If urine were collected in 2-hr. aliquots, the max. sp. activity occurred about 2 hrs. earlier in all gouty subjects than in controls, and the peak values for the 3 hyperexcretors were 2-4-fold greater than controls. There was an increased I turnover in the 3 hyperexcretor gouty subjects, but there may be a continuous gradation in the magnitude of purine synthesis in man.

IT **2888-19-9, Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-phosphate
(in urine in gout)

RN 2888-19-9 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 18 OF 26 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1962:465523 CAPLUS

DOCUMENT NUMBER: 57:65523

ORIGINAL REFERENCE NO.: 57:13071h-i

TITLE: Enzymic synthesis of a riboside involved in histamine metabolism

AUTHOR(S): Fernandes, J. F.; Castellani, Olga; Plese, Mitzi

CORPORATE SOURCE: Univ. Sao Paulo Med. School, Brazil

SOURCE: Ciencia Cult. (Sao Paulo) (1961), 13(No. 2), 87-92

DOCUMENT TYPE: Journal

LANGUAGE: Unavailable

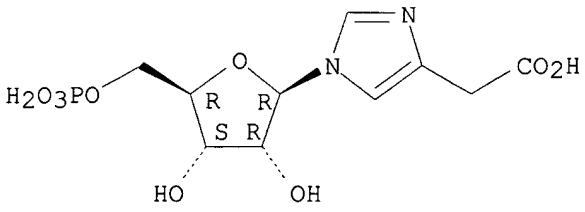
AB Exts. from the small intestine mucosa of the dog catalyzed reactions between histamine and **imidazole** acetate with 5-phosphoribosyl pyrophosphate. Exts. from the ileum and from the lung of the guinea pig also catalyzed these reactions, which required the presence of adenosine phosphate and of phosphate ions. A method for the purification and characterization of the products is presented.

IT **2888-19-9, Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-phosphate
(formation by intestinal enzymes and its isolation and characterization)

RN 2888-19-9 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 19 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1962:79591 CAPLUS
DOCUMENT NUMBER: 56:79591
ORIGINAL REFERENCE NO.: 56:15596b-h
TITLE: Synthesis of 1-(.beta.-D-ribofuranosyl) **imidazole**-4(or 5)-acetonitrile, 1-(.beta.-D-ribofuranosyl)**imidazole**-4(or 5)-acetic acid, and 4(or 5)-(2-aminoethyl)-1-(.beta.-D-ribofuranosyl)**imidazole**
AUTHOR(S): Bauer, Hugo
CORPORATE SOURCE: Natl. Insts. of Health, Bethesda, MD
SOURCE: J. Org. Chem. (1962), 27, 167-70
CODEN: JOCEAH; ISSN: 0022-3263
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable

AB cf. CA 53, 6214a. Condensation of the HgCl2 complex (I) of **imidazoleacetonitrile** (II) with 2,3,5-tri-O-benzoyl-D-ribosyl bromide (III) and subsequent debenzoylation gave 1-(.beta.-D-ribofuranosyl)**imidazole**-4(or 5)acetonitrile (IV), converted by hydrolysis or catalytic hydrogenation to the corresponding acid (V) and histamine ribose (VI). II (2.14 g.) and 2.12 g. Na2CO3 in 150 ml. hot H2O contg. 3 g. Celite stirred with addn. of 5.43 g. HgCl2 in 150 ml. hot H2O, the centrifuged ppt. washed twice with H2O, dried at 70.degree. in vacuo, freed from moisture by azeotropic distn. with xylene, suspended in 150 ml. xylene, the suspension gently refluxed 3 hrs. with stirring with III (prepd. from 9 g. 2,3,5-tri-O-benzoyl-.beta.-D-ribose) in 100 ml. xylene, the xylene evapd. from the filtered soln., the light brown oil (VII) washed with Et2O, the insol. residue (5 g.) taken up in hot alc., the cooled soln. decanted, and the chilled liquid dild. with petr. ether yielded 1.6 g. HgCl2 complex, (C31H25N3O7)2.HgCl2, m. 95.degree.. VII in CH2Cl2 shaken with 30% aq. KI, the CH2Cl2 soln. evapd., the residual oil taken up in alc., pptd. by addn. of petr. ether, and the middle fraction dried in vacuo at 45.degree. yielded 1-(2,3,5-tri-O-benzoyl-.beta.-D-ribosyl)-**imidazole**-4(or 5)-acetonitrile HCl salt (VIII), softening at 90.degree.. VII (10 g.) shaken in CH2Cl2 with 30% aq. KI, the dried CH2Cl2 soln. evapd., the oil taken up in 25 ml. MeOH, kept 16 hrs. at 20.degree. with 10 ml. 2N (MeO)2Ba, the soln. adjusted to Congo red with 2N H2SO4, centrifuged, the supernatant washed with Et2O, freed from volatile materials, the soln. (IX) adsorbed on Dowex 50, eluted with 1 l. 2N H2SO4, the eluate treated with Ba(OH)2, the neutral soln. made slightly alk. with 10% NH4OH, evapd. at 30.degree. in vacuo, the residue taken up in H2O, decolorized (Norit) at room temp., evapd., the colorless gum (1.7 g.) taken up in abs. alc., the filtered soln. evapd., the product purified by soln. in alc. and filtration of the cooled soln., the filtrate evapd. in vacuo, washed with Me2CO, and dried gave hydrated IV, C10H13N3O4.H2O, with slight hydrolysis of the CN group. IV reacted with 1 equiv. NaIO4 on titration showing the presence of 1 mole ribose. IX boiled 2 hrs. with 15 g. Ba(OH)2 in 150 ml. H2O with evolution of NH3, treated with 2N H2SO4, the decolorized filtrate chromatographed on Dowex 1 acetate, eluted gradually with 3N AcOH to give material contg. free **imidazoleacetic** acid, the following fractions acidified with 2N

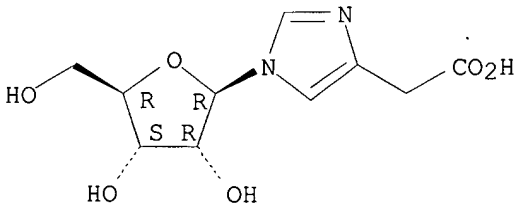
HCl, the soln. evapd., and the residue (1.5 g.) recrystd. from H2O-Me2CO yielded VI HCl salt, m. 135%, [.alpha.]20D -37.degree. (c 1.0, H2O), -51.4.degree. (c 1.16, MeOH), infrared spectrum in Nujol identical with that of a compd. isolated from the urine of rats injected with histamine or **imidazoleacetic** acid (CA 49, 11135b). II (1.07 g.) in 20 ml. alc. and 2 ml. concd. H2SO4 hydrogenated 30 hrs. with shaking with 0.1 g. prereduced PtO2, the mixt. adsorbed on Dowex 50, and eluted with 4N HCl gave 1.0 g. histamine di-HCl salt. VII (50 ml.) acidified with 0.5 ml. concd. HC2SO4, hydrogenated with 0.3 g. prereduced PtO2 added in 2 portions, the acid removed as BaSO4, the soln. percolated through Dowex 50 H+, the column eluted with 2N HCl, and the product crystd. from H2O-Et2O gave VI di-HCl salt, m. 174-5.degree..

IT **95445-10-6**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-, hydrochloride
(prepn. of)

RN 95445-10-6 CAPLUS

CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl-, hydrochloride (6CI, 7CI) (CA INDEX NAME)

Absolute stereochemistry.



● HCl

L10 ANSWER 20 OF 26 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1962:74743 CAPLUS

DOCUMENT NUMBER: 56:74743

ORIGINAL REFERENCE NO.: 56:14584b-d

TITLE: The catabolism of tissue nucleic acid. III. The catabolism of ribonucleic acid after total-body x-irradiation

AUTHOR(S): Gerber, Georg B.; Gerber, Gisela; Altman, Kurt I.

CORPORATE SOURCE: Univ. of Rochester, Rochester, NY

SOURCE: Intern. J. Radiation Biol. (1961), 4, 67-73

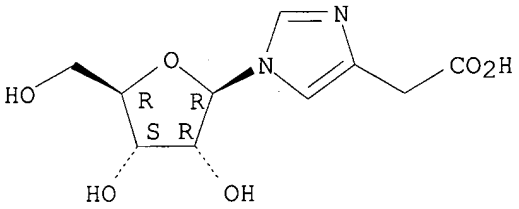
DOCUMENT TYPE: Journal

LANGUAGE: Unavailable

AB cf. CA 54, 25159h. The effect of total-body x-irradiation on ribonucleic acid (RNA) catabolism was studied in rats whose RNA had been labeled by injection of glucose-U-C14 three days previously. The sp. activity of urinary ribosyl **imidazole** acetate (I) as well as of RNA of liver, intestine, muscle, spleen and thymus was detd. after x- or sham-irradiation. Rats were either pair fed or starved after irradiation. After 1000 r. the sp. activity of I was increased whereas that of intestinal and muscle RNA was decreased with little change in liver RNA. In pair-fed animals, exposure to 756 r. decreased the sp. activity of spleen and thymus RNA, and gave a steady decrease in that of I over 5 days. Sp. activity of liver RNA was diminished in the sham-irradiated rats and was equiv. to that of I which was excreted on the first day after treatment. It was concluded that radiation-induced increase in RNA catabolism is present mainly in intestine and muscle on the second and third day after exposure whereas starvation-induced increase in catabolism occurs primarily in the liver and on the first day of starvation.

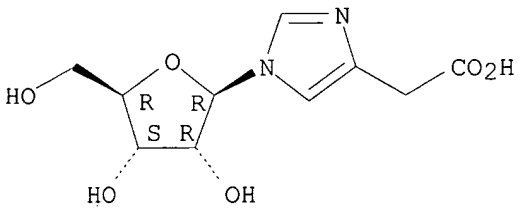
IT **29605-99-0, Imidazole**-4-acetic acid, 1-ribosyl-
(in urine after x-ray irradiation)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 21 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1960:120119 CAPLUS
DOCUMENT NUMBER: 54:120119
ORIGINAL REFERENCE NO.: 54:23006e-g
TITLE: Histamine metabolism in human disease
AUTHOR(S): Beall, Gildon N.; Van Arsdel, Paul P., Jr.
CORPORATE SOURCE: Univ. of Washington, Seattle
SOURCE: J. Clin. Invest. (1960), 39, 676-83
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable
AB The metabolism of histamine-C14 was studied in 7 control subjects, 5 patients with Laennec's cirrhosis, 3 with bronchial asthma, 3 with histaminic cephalgia, and 1 with uremia due to chronic glomerulonephritis. Radioactive histamine was always rapidly cleared from the blood, and its excretion was usually complete in 30 hrs. Sepn. of the excreted radioactive metabolites by paper chromatography and radioautography showed **imidazoleacetic** acid to be the principal product of histamine metabolism. **Imidazoleacetic** acid riboside and 1,4-methylimidazoleacetic acid were also consistently recovered in the urine, but histamine-C14 was not detected. Histamine inactivation and degradation in man is a rapid and complete process and, by the methods employed, no abnormalities were found in the diseases studied. 25 references.
IT **29605-99-0, Imidazole**-4-acetic acid, 1-ribosyl-
(in urine as histamine metabolite)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 22 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1959:28685 CAPLUS
DOCUMENT NUMBER: 53:28685
ORIGINAL REFERENCE NO.: 53:5140c-e
TITLE: Synthesis of 1-.beta.-D-ribofuranosyl-4(5)-

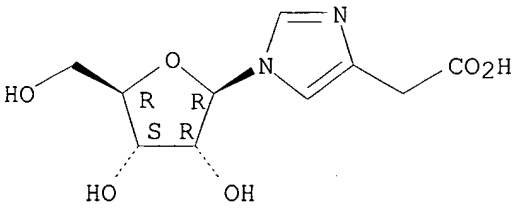
glyoxalinylacetic acid, a metabolite of histamine
AUTHOR(S): Baddiley, J.; Buchanan, J. G.; Hayes, D. H.; Smith, P. A.
CORPORATE SOURCE: Univ. Durham, Newcastle-upon-Tyne, UK
SOURCE: J. Chem. Soc. (1958) 3743-5
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable

AB A synthesis is described which established the configuration of the ribosyl linkage: the HgCl2 salt of Me 4-glyoxalinyl acetate is condensed with tri-O-benzoyl-.beta.-D-ribofuranosyl chloride in boiling xylene, the product debenzoylated, and the Me ester group hydrolyzed. The ribosyl compd. is isolated by ion-exchange chromatography. Comparison with the natural compd. (m.p., infrared spectrum, RF values in various solvent systems, and behavior on acid hydrolysis) proves their identity. Reaction of a tri-O-acetyl-or tri-O-benzoylribofuranosyl halide with Hg salts yields only the .beta.-anomer in all cases so far studied. This established the .beta.-configuration of the natural product.

IT **29605-99-0**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-(?) **95445-10-6**, **Imidazole**-4-acetic acid, 1-.beta.-D-ribofuranosyl-(?), hydrochloride
(prepn. of)

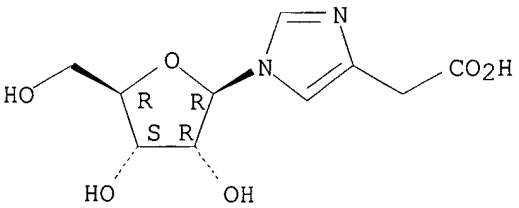
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 95445-10-6 CAPLUS
CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl-, hydrochloride (6CI, 7CI) (CA INDEX NAME)

Absolute stereochemistry.

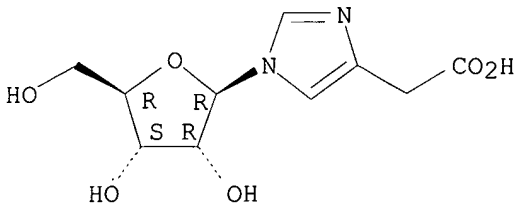


● HCl

L10 ANSWER 23 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1959:24133 CAPLUS
DOCUMENT NUMBER: 53:24133
ORIGINAL REFERENCE NO.: 53:4472f
TITLE: Isolation of **imidazoleacetic** acid riboside
AUTHOR(S): Tabor, H.
CORPORATE SOURCE: Natl. Inst. of Arthritis and Metabolic Diseases,

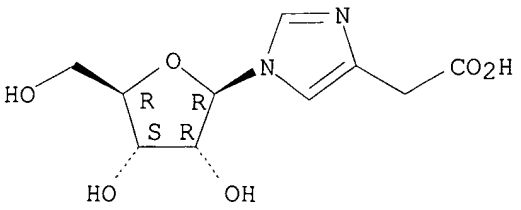
Bethesda, MD
SOURCE: Ciba Foundation Symposium, Histamine (1956) 51
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable
AB Reviews with many references.
IT **29605-99-0, Imidazoleacetic** acid, 1-ribosyl-
(isolation of)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 24 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1959:12622 CAPLUS
DOCUMENT NUMBER: 53:12622
ORIGINAL REFERENCE NO.: 53:2404a-c
TITLE: Ribose metabolism. V. Factors influencing in vivo ribose synthesis in the rat
AUTHOR(S): Hiatt, Howard H.
CORPORATE SOURCE: Harvard Med. School, Boston, MA
SOURCE: J. Clin. Invest. (1958), 37, 1453-60
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable
AB cf. C.A. 52, 16519c. Ribose synthesis in vivo was studied by isolating **imidazoleacetic** acid (IZA) riboside from rats given IZA and a C14-labeled sugar. Evidence is presented which indicates an impairment of riboside excretion in partially hepatectomized animals and in diabetic animals. The isotope distribution in ribose synthesized from glucose-2-C14 by normal animals is consistent with synthesis via both the oxidative and the nonoxidative reactions of the pentose phosphate pathway. Thiamine deficiency resulted in a marked decrease of ribose synthesis from hexose via the nonoxidative mechanism. An apparent increase in ribose production by way of the oxidative reactions was observed in rats with regenerating livers and in tumor-bearing rats. Evidence is presented for the direct incorporation of administered ribose into the urinary riboside.
IT **29605-99-0, Imidazoleacetic** acid, 1-ribosyl-
(in urine as ribose metabolite)
RN 29605-99-0 CAPLUS
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 25 OF 26 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1957:52995 CAPLUS
DOCUMENT NUMBER: 51:52995
ORIGINAL REFERENCE NO.: 51:9851f-h
TITLE: The metabolism of histamine in various species
AUTHOR(S): Schayer, Richard W.
CORPORATE SOURCE: Rheumatic Fever Research Inst., Chicago
SOURCE: Brit. J. Pharmacol. (1956), 11, 472-3
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable

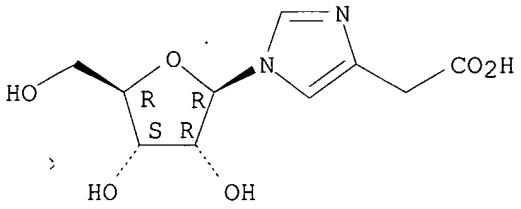
AB cf. C.A. 50, 14838a. Quant. analyses for histamine (I) metabolites in urine of various species after feeding or injecting I-C14 were performed. In the rabbit, I was equally metabolized by oxidation and by methylation to give 1-methylimidazole-4-acetic acid (II) and 1-ribosylimidazole-4(5)-acetic acid (III). The mouse metabolized fed I to III plus some **imidazole**-4-acetic acid and II. In cats and man methylation was the principal route of metabolism for fed or injected I to give II. The cat excreted some unchanged I. The dog excreted injected I as II together with 1-methyl-4-(2-aminoethyl)**imidazole** and 4-**imidazoleacetic** acid. In the cat a small fraction of injected L-histidine-C14 was recovered as tissue I after 8 days; in contrast no injected I-C14 was retained.

IT **29605-99-0, Imidazole**-4-acetic acid, 1-ribosyl-
(in histidine metabolism by various species)

RN 29605-99-0 CAPLUS

CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

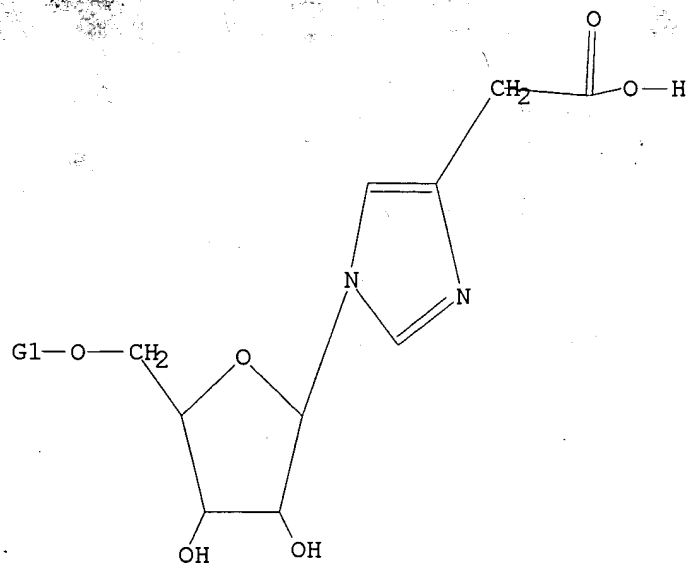


L10 ANSWER 26 OF 26 MEDLINE
ACCESSION NUMBER: 95341318 MEDLINE
DOCUMENT NUMBER: 95341318 PubMed ID: 7616240
TITLE: **Imidazoleacetic** acid, a gamma-aminobutyric acid receptor agonist, can be formed in rat brain by oxidation of histamine.
AUTHOR: Thomas B; Prell G D
CORPORATE SOURCE: Department of Pharmacology, Mount Sinai School of Medicine, City University of New York, New York, USA.
CONTRACT NUMBER: NS 28012 (NINDS)
SOURCE: JOURNAL OF NEUROCHEMISTRY, (1995 Aug) 65 (2) 818-26.
Journal code: 2985190R. ISSN: 0022-3042.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 199508
ENTRY DATE: Entered STN: 19950905
Last Updated on STN: 19980206
Entered Medline: 19950824

AB It is generally accepted that in mammalian brain histamine is metabolized solely by histamine methyltransferase (HMT), to form tele-methylhistamine, then oxidized to tele-methylimidazoleacetic acid. However, histamine's oxidative metabolite in the periphery, **imidazoleacetic** acid

(IAA), is also present in brain and CSF, and its levels in brain increase after inhibition of HMT. To reinvestigate if brain has the capacity to oxidize histamine and form IAA, conscious rats were injected with [3H]histamine (10 ng), either into the lateral ventricles or cisterna magna, and decapitated 30 min later. In brains of saline-treated rats, most radioactivity recovered was due to tele-methylhistamine and tele-methylimidazoleacetic acid. However, significant amounts of tritiated IAA and its metabolites, IAA-ribotide and IAA-riboside, were consistently recovered. In rats pretreated with metoprine, an inhibitor of HMT, labeled IAA and its metabolites usually comprised the majority of histamine's tritiated metabolites. [3H]Histamine given intracisternally produced only trace amounts of oxidative metabolites. Formation of IAA, a potent GABA-A agonist with numerous neurochemical and behavioral effects, from minute quantities of histamine in brain indicates a need for reevaluation of histamine's metabolic pathway or pathways in brain and suggests a novel mechanism for interactions between histamine and the GABAergic system.

L1 STR



G1 PO3H2,H

Structure attributes must be viewed using STN Express query preparation.

=> sL1 sss full
SL1 IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).

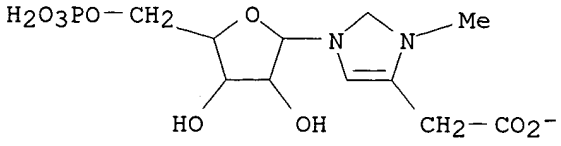
=> s L1 sss full
FULL SEARCH INITIATED 18:45:34 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 197 TO ITERATE

100.0% PROCESSED 197 ITERATIONS 5 ANSWERS
SEARCH TIME: 00.00.01

L2 5 SEA SSS FUL L1

=> d L2 1-5

L2 ANSWER 1 OF 5 REGISTRY COPYRIGHT 2001 ACS
RN 142606-76-6 REGISTRY
CN 1H-Imidazolium, 4-(carboxymethyl)-3-methyl-1-(5-O-phosphono-.beta.-D-ribofuranosyl)-, inner salt, monosodium salt (9CI) (CA INDEX NAME)
MF C11 H17 N2 O9 P . Na
SR CA
LC STN Files: CA, CAPLUS

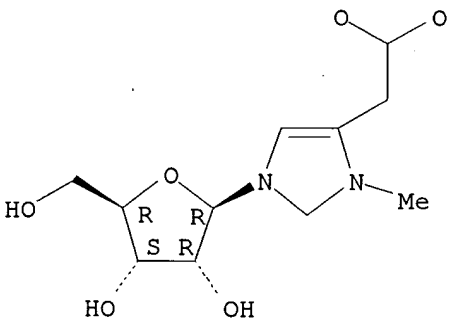


● Na

*** FRAGMENT DIAGRAM IS INCOMPLETE ***
1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

L2 ANSWER 2 OF 5 REGISTRY COPYRIGHT 2001 ACS
RN 142527-55-7 REGISTRY
CN 1H-Imidazolium, 4-(carboxymethyl)-3-methyl-1-.beta.-D-ribofuranosyl-, inner salt (9CI) (CA INDEX NAME)
FS STEREOSEARCH
MF C11 H16 N2 O6
SR CA
LC STN Files: CA, CAPLUS

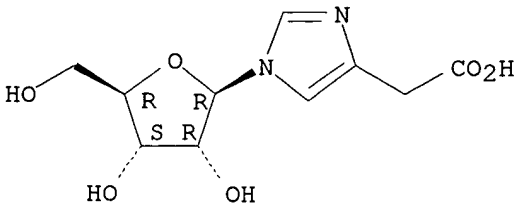
Absolute stereochemistry.



*** FRAGMENT DIAGRAM IS INCOMPLETE ***
1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

L2 ANSWER 3 OF 5 REGISTRY COPYRIGHT 2001 ACS
RN 95445-10-6 REGISTRY
CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl-, hydrochloride (6CI, 7CI) (CA INDEX NAME)
FS STEREOSEARCH
MF C10 H14 N2 O6 . Cl H
LC STN Files: BEILSTEIN*, CAOLD
(*File contains numerically searchable property data)
CRN (29605-99-0)

Absolute stereochemistry.

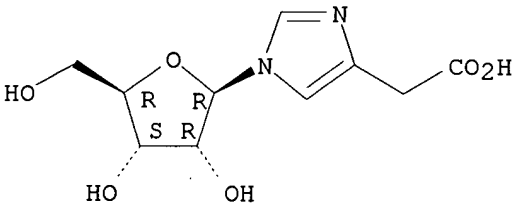


● HCl

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L2 ANSWER 4 OF 5 REGISTRY COPYRIGHT 2001 ACS
RN 29605-99-0 REGISTRY
CN 1H-Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl- (6CI, 7CI, 8CI)
OTHER NAMES:
CN 1-Ribosylimidazole-4-acetic acid
CN Imidazoleacetic acid riboside
FS STEREOSEARCH
DR 55348-45-3
MF C10 H14 N2 O6
CI COM
LC STN Files: BEILSTEIN*, BIOSIS, CA, CAOLD, CAPLUS, MEDLINE, TOXLIT
(*File contains numerically searchable property data)

Absolute stereochemistry.

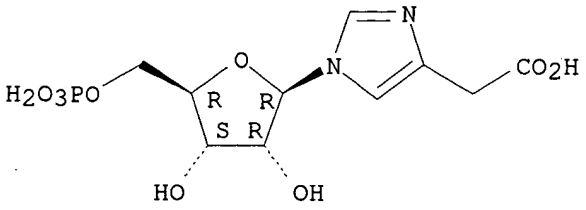


14 REFERENCES IN FILE CA (1967 TO DATE)
14 REFERENCES IN FILE CAPLUS (1967 TO DATE)
10 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L2 ANSWER 5 OF 5 REGISTRY COPYRIGHT 2001 ACS
RN 2888-19-9 REGISTRY
CN 1H-Imidazole-4-acetic acid, 1-(5-O-phosphono-.beta.-D-ribofuranosyl)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-(dihydrogen phosphate) (8CI)
CN Imidazole-4-acetic acid, 1-.beta.-D-ribofuranosyl-, 5'-phosphate (6CI, 7CI)
OTHER NAMES:
CN 1-Ribosylimidazole-4-acetic acid 5'-phosphate
CN Imidazole-4-acetic acid ribotide

FS STEREOSEARCH
MF C10 H15 N2 O9 P
LC STN Files: CA, CAOLD, CAPLUS, TOXLIT

Absolute stereochemistry.



2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)
6 REFERENCES IN FILE CAOLD (PRIOR TO 1967)